

PALMA HAND SANITIZER- ethyl alcohol gel
UNITRADE FZE

Disclaimer: Most OTC drugs are not reviewed and approved by FDA, however they may be marketed if they comply with applicable regulations and policies. FDA has not evaluated whether this product complies.

PALMA DE SALUS HAND SANITIZER

Drug Facts

Active ingredient

Ethyl alcohol 70% v/v

Purpose

Antiseptic

Uses

• hand sanitizer to decrease bacteria on the skin • recommended for repeated use • for use when soap and water are not available

Warnings

Flammable, keep away from fire/flame
For external use only

Do not use • in children less than 2 months of age • on open skin wounds

When using this product • do not get into eyes. In case of contact, rinse eyes thoroughly with water

Stop use and ask a doctor if • irritation and redness develop • condition persists for more than 72 hours

Keep out of reach of children. If swallowed, get medical help or contact a Poison Control Center right away.

Directions

• wet hands thoroughly with product and allow to dry without wiping • supervise children under 6 years of age when using this product to avoid swallowing

Other information

• store between 15-30°C (59-86°F) • avoid freezing and excessive heat above 40°C (104°F)

Inactive ingredients

water, caprylic /capric triglyceride, avocado (*Persea gratissima*) oil, propylene glycol, glycerin, jojoba (*Simmondsia chinensis*) seed oil, peg-40 hydrogenated castor oil, polyacrylamide, c13-14 isoparaffin, laureth-7, acrylates / c10-30 alkyl acrylates crosspolymer, niacinamide, sodium hyaluronate, tocopheryl acetate, panthenol, gotu kola (*Centella asiatica*) root extract, triethanolamine, fragrance.

Questions? 1-516-535-5555

You may also report serious side effects to this phone number. Mon-Fri 9:00 AM - 5:00 PM

MOISTURIZING

WITH AVOCADO OIL, JOJOBA OIL, VITAMIN B3 & B5

PROTECT. HYDRATE. NOURISH.

Distributed by:
Palma De Salus, Inc
320 Country Road, Suite 205
Garden City, NY 11530, U.S.A

Made in U.A.E.
Designed in Los Angeles
palmadesalus.com

Packaging

PALMA HAND SANITIZER				
ethyl alcohol gel				
Product Information				
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:81286-101	
Route of Administration	TOPICAL			
Active Ingredient/Active Moiety				
Ingredient Name		Basis of Strength	Strength	
ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)		ALCOHOL	70 mL in 100 mL	
Inactive Ingredients				
Ingredient Name			Strength	
WATER (UNII: 059QF0K00R)				
MEDIUM-CHAIN TRIGLYCERIDES (UNII: C9H2L21V7U)				
AVOCADO OIL (UNII: 6VNO72PFC1)				
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)				
GLYCERIN (UNII: PDC6A3C0OX)				
JOJOBA OIL (UNII: 724GKU717M)				
POLYOXYL 40 HYDROGENATED CASTOR OIL (UNII: 7YC686GQ8F)				
POLYACRYLAMIDE (1300000 MW) (UNII: SC5Y4X78TG)				
C13-14 ISOPARAFFIN (UNII: E4F12ROE70)				
LAURETH-7 (UNII: Z95S6G8201)				
CARBOMER INTERPOLYMER TYPE A (ALLYL SUCROSE CROSSLINKED) (UNII: 59TL3WG5CO)				
NIACINAMIDE (UNII: 25X51I8RD4)				
HYALURONATE SODIUM (UNII: YSE9PPT4TH)				
.ALPHA.-TOCOPHEROL ACETATE (UNII: 9E8X80D2L0)				
PANTHENOL (UNII: WV9CM0O67Z)				
CENTELLA ASIATICA LEAF (UNII: 6810070TYD)				
TROLAMINE (UNII: 9O3K93S3TK)				
Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:81286-101-10	1 in 1 CARTON	07/26/2021	
1		325 mL in 1 BOTTLE, PUMP; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333A	07/26/2021	

Labeler - UNITRADE FZE (864268417)

Revised: 7/2021

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